

NCI Trial ID: NCI-2016-01182 ▾

Trial Overview ▾

Trial Identification

Trial History

Trial Milestones

On Hold Info

View TSR

Assign Ownership

Check Out History

Email Logs

Administrative Data ▾

General Trial Details

Regulatory Information - FDAAA

Regulatory Information - Human Subject Safety

Regulatory Information - IND/IDE

Trial Status

Trial Funding

Participating Sites

Collaborators

Trial Related Documents

NCI Specific Information

Scientific Data ▾

Trial Design

Trial Description

Interventions

Arms/Groups

Eligibility Criteria

Associated Trials

Diseases/Conditions

Data Table 4 Anatomic Sites

Outcome Measures

Sub-Groups Stratification

Biomarkers

Completion ▾

Abstraction Validation

Results Information

≡ Trial Overview (NCI Trial ID: NCI-2016-01182) ⌵

QA - Team Title - Dolor qui et nesciunt, ea culpa illo saepe incididunt sapiente enim proident, et similique veritatis quis mollitia et repudiandae assumenda.

NCI ID: NCI-2016-01182

Principal Investigator: James Lee

Information Source: Protocol

NCT ID:

Clinical Research Category: Interventional

Data Table 4 Funding Source: Memorial Hospital Colorado Springs

Lead Organization Trial ID: QA - Team Dolor qui et nesciun

Last Submitter: ctrptrialsubmitter3

Current Trial Status: In Review

Lead Organization: Harborview Medical Center

Last Submitter Organization: ZZZ test org for test accounts

Current Trial Status Date: 06-Jul-2016

Submission Method: Registry

Last Updated By: ctrpabstractor3

Processing Status: Accepted

Amendment Number:

Last Updated Date: 29-Jul-2016 10:15

Checked Out for Admin. Use by: ctrpabstractor3

Amendment Date:

Checked Out for Scientific. Use by: ctrpabstractor3

[← Back to Search Results](#)

[Admin Check In](#)

[Scientific Check In](#)

[Admin/Scientific Check In](#)

List of Participating Sites

[Add Participating Site](#)

NCI Trial ID: NCI-2016-01182 ▾

Trial Overview ▾

Trial Identification

Trial History

Trial Milestones

On Hold Info

View TSR

Assign Ownership

Check Out History

Email Logs

Administrative Data ▾

General Trial Details

Regulatory Information - FDAAA

Regulatory Information - Human Subject Safety

Regulatory Information - IND/IDE

Trial Status

Trial Funding

Participating Sites

Collaborators

Trial Related Documents

NCI Specific Information

Scientific Data ▾

Trial Design

Trial Description

Interventions

Arms/Groups

Eligibility Criteria

Associated Trials

Diseases/Conditions

Data Table 4 Anatomic Sites

Outcome Measures

Sub-Groups Stratification

Biomarkers

Completion ▾

Abstraction Validation

Results Information

Trial Overview (NCI Trial ID: NCI-2016-01182) ▾

QA - Team Title - Dolor qui et nesciunt, ea culpa illo saepe incididunt sapiente enim proident, et similique veritatis quis mollitia et repudiandae assumenda.

NCI ID: NCI-2016-01182

Principal Investigator: James Lee

Information Source: Protocol

NCIT ID:

Clinical Research Category: Interventional

Data Table 4 Funding Source: Memorial Hospital Colorado Springs

Lead Organization Trial ID: QA - Team Dolor qui et nesciunt

Last Submitter: ctriptrialsubmitter3

Current Trial Status: In Review

Lead Organization: Harborview Medical Center

Last Submitter Organization: ZZZ test org for test accounts

Current Trial Status Date: 06-Jul-2016

Submission Method: Registry

Last Updated By: ctripabstractor3

Processing Status: Accepted

Amendment Number:

Last Updated Date: 29-Jul-2016 10:15

Checked Out for Admin. Use by: ctripabstractor3

Amendment Date:

Checked Out for Scientific. Use by: ctripabstractor3

← Back to Search Results

Admin Check In

Scientific Check In

Admin/Scientific Check In

Abstraction Validation

Abstraction Validation Failed: Errors in Administrative Data: 3

Description	Comment
Review Board Approval Status is null	Select [Regulatory Information - Human Subject Safety] from Administrative Data menu.
duplicate sites	[Select Participating Sites] from Administrative Data menu.
duplicate Investigators at the same site	[Select Participating Sites] from Administrative Data menu.

Abstraction Validation

Description	Comment
Clinical Research Category=Interventional AND Masking is null	[Select Trial Design] from Scientific Data menu.
Clinical Research Category = 'Interventional' AND Intervention Model is null	[Select Trial Design] from Scientific Data menu.
Number of Arms / Groups is null	[Select Trial Design] from Scientific Data menu.
Clinical Research Category = 'Interventional' AND Allocations is null	[Select Trial Description] from Scientific Data menu.
Brief Title is null	[Select Trial Description] from Scientific Data menu.
Brief Title is duplicate	[Select Trial Description] from Scientific Data menu.
Brief Summary is null	[Select Trial Description] from Scientific Data menu.
Eligibility Criteria is null	[Select Eligibility Criteria] from Scientific Data menu.
Accepts Healthy Volunteers on Eligibility is null	[Select Eligibility Criteria] from Scientific Data menu.
Gender is null	[Select Eligibility Criteria] from Scientific Data menu.
Minimum Age and	[Select Eligibility Criteria] from Scientific Data menu.
Maximum Age and	[Select Eligibility Criteria] from Scientific Data menu.
Other Criteria is null	[Select Eligibility Criteria] from Scientific Data menu.
Disease/Condition is null	[Select Diseases/Conditions] from Scientific Data menu.
Primary Outcome is null	[Select Outcome Measures] from Scientific Data menu.

Abstraction Validation Warnings: 12

Description	Comment
Current Trial Status = IN REVIEW and Board Approval Status is not Submitted; Pending	[Select Regulatory Information Human Subject Safety] from Administrative Data menu.
Trial Status is In Reviewand Board Approval Status is not Submitted; Pending	[Select Regulatory Information Human Subject Safety] from Administrative Data menu.
Study Status = APPROVED and Site Status = ENROLLING BY INVITATION	[Select Participating Sites] from Administrative Data menu.
Study Status = IN REVIEW and Site Status = ENROLLING BY INVITATION	[Select Participating Sites] from Administrative Data menu.
Study Status WITH	[Select Participating Sites] from Administrative Data menu.
Participating Site =	[Select Participating Sites] from Administrative Data menu.
Primary Site status has been set to Nullified	[Select Participating Sites] from Administrative Data menu.
Primary Investigator has been set to Nullified	[Select Participating Sites] from Administrative Data menu.
Primary Contact status has been set to Nullified	[Select Participating Sites] from Administrative Data menu.
Central Contact is null AND no primary contact information (person; phone OR email) for each participating site	[Select Participating Sites] from Administrative Data menu.
Target accrual is null	[Select Participating Sites] from Administrative Data menu.
Arm is null	[Select Arms/Groups] from Scientific Data menu.